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U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE

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TRANSMITTAL LETTER TO THE UNITED STATES  
DESIGNATED/ELECTED OFFICE (DO/EO/US)  
CONCERNING A FILING UNDER 35 U.S.C. 371

40740

U.S. APPLICATION NO. (If known, see 37 CFR 1.5)

09/646835

INTERNATIONAL APPLICATION NO.  
PCT/EP99/02165INTERNATIONAL FILING DATE  
29 March 1999PRIORITY DATE CLAIMED  
27 March 1998TITLE OF INVENTION  
NEW USE OF Hsp70 PROTEINAPPLICANT(S) FOR DO/EO/US  
Gabriele MÜLTHOFF

Applicant herewith submits to the United States Designated/Elected Office (DO/EO/US) the following items and other information:

1. ☒ This is a **FIRST** submission of items concerning a filing under 35 U.S.C. 371.
2. ☐ This is a **SECOND** or **SUBSEQUENT** submission of items concerning a filing under 35 U.S.C. 371.
3. ☐ This express request to begin national examination procedures (35 U.S.C. 371(f)) at any time rather than delay examination until the expiration of the applicable time limit set in 35 U.S.C. 371(b) and PCT Articles 22 and 39(1).
4. ☒ A proper Demand for International Preliminary Examination was made by the 19th month from the earliest claimed priority date.
5. ☒ A copy of the International Application as filed (35 U.S.C. 371(c)(2))
  - a. ☒ is transmitted herewith (required only if not transmitted by the International Bureau).
  - b. ☐ has been transmitted by the International Bureau.
  - c. ☐ is not required, as the application was filed in the United States Receiving Office (RO/US).
6. ☒ A translation of the International Application into English (35 U.S.C. 371(c)(2)).
7. ☒ Amendments to the claims of the International Application under PCT Article 19 (35 U.S.C. 371(c)(3))
  - a. ☐ are transmitted herewith (required only if not transmitted by the International Bureau).
  - b. ☒ have been transmitted by the International Bureau.
  - c. ☐ have not been made; however, the time limit for making such amendments has NOT expired.
  - d. ☐ have not been made and will not be made.
8. ☒ A translation of the amendments to the claims under PCT Article 19 (35 U.S.C. 371(c)(3)).
9. ☐ An oath or declaration of the inventor(s) (35 U.S.C. 371(c)(4)).
10. ☐ A translation of the annexes to the International Preliminary Examination Report under PCT Article 36 (35 U.S.C. 371(c)(5)).

**Items 11. to 16. below concern document(s) or information included:**

11. ☒ An Information Disclosure Statement under 37 CFR 1.97 and 1.98.
12. ☐ An assignment document for recording. A separate cover sheet in compliance with 37 CFR 3.28 and 3.31 is included.
13. ☐ A FIRST preliminary amendment.  
☐ A SECOND or SUBSEQUENT preliminary amendment.
14. ☐ A substitute specification.
15. ☐ A change of power of attorney and/or address letter.
16. ☒ Other items or information:
  - International Search Report
  - International Preliminary Examination Report
  - Request
  - Written Opinion
  - Response to Written Opinion of 15 May 2000
  - Communication of 3 July 2000
  - Response Thereto of 12 July 2000

09/646835

17. ☒ The following fees are submitted:**BASIC NATIONAL FEE (37 CFR 1.492 (a) (1) - (5)) :**

Neither international preliminary examination fee (37 CFR 1.482)  
nor international search fee (37 CFR 1.445(a)(2)) paid to USPTO  
and International Search Report not prepared by the EPO or JPO ..... \$970.00

International preliminary examination fee (37 CFR 1.482) not paid to  
USPTO but International Search Report prepared by the EPO or JPO ..... \$840.00

International preliminary examination fee (37 CFR 1.482) not paid to USPTO but  
international search fee (37 CFR 1.445(a)(2)) paid to USPTO ..... \$690.00

International preliminary examination fee paid to USPTO (37 CFR 1.482)  
but all claims did not satisfy provisions of PCT Article 33(1)-(4) ..... \$670.00

International preliminary examination fee paid to USPTO (37 CFR 1.482)  
and all claims satisfied provisions of PCT Article 33(1)-(4) ..... \$96.00

**ENTER APPROPRIATE BASIC FEE AMOUNT =****CALCULATIONS** PTO USE ONLY

422 Rec'd PCT/PTO 22 SEP 2000

\$ 840.00

Surcharge of **\$130.00** for furnishing the oath or declaration later than ☐ 20 ☐ 30  
months from the earliest claimed priority date (37 CFR 1.492(e)).

\$

CLAIMS	NUMBER FILED	NUMBER EXTRA	RATE
Total claims	34 - 20 =	14	X \$18.00
Independent claims	4 - 3 =	1	X \$78.00
MULTIPLE DEPENDENT CLAIM(S) (if applicable)			+ \$260.00

\$ 252.00

\$ 78.00

\$ 260.00

**TOTAL OF ABOVE CALCULATIONS =**

\$ 1430.00

Reduction of 1/2 for filing by small entity, if applicable. A Small Entity Statement  
must also be filed (Note 37 CFR 1.9, 1.27, 1.28).

\$

**SUBTOTAL =**

\$ 715.00

Processing fee of **\$130.00** for furnishing the English translation later than ☐ 20 ☐ 30  
months from the earliest claimed priority date (37 CFR 1.492(f)).

\$ 715.00

**TOTAL NATIONAL FEE =**

\$ 715.00

Fee for recording the enclosed assignment (37 CFR 1.21(h)). The assignment must be  
accompanied by an appropriate cover sheet (37 CFR 3.28, 3.31). **\$40.00** per property

\$

**TOTAL FEES ENCLOSED =**

\$ 715.00

Amount to be  
**refunded:** \$  
**charged:** \$

a. ☒ A check in the amount of \$ 715.00 to cover the above fees is enclosed.

b. ☐ Please charge my Deposit Account No. \_\_\_\_\_ in the amount of \$ \_\_\_\_\_ to cover the above fees.  
A duplicate copy of this sheet is enclosed.

c. ☒ The Commissioner is hereby authorized to charge any additional fees which may be required, or credit any  
overpayment to Deposit Account No. 18-2220. A duplicate copy of this sheet is enclosed.

The Declaration/Power of Attorney, Assignment and Small Entity Declaration will be filed shortly.

Priority is claimed from DE Ser. No. 198 13 760.5 filed 27 March 1998 and EP99/02056 filed 26 March 1999.

**NOTE: Where an appropriate time limit under 37 CFR 1.494 or 1.495 has not been met, a petition to revive (37 CFR 1.137(a) or (b)) must be filed and granted to restore the application to pending status.**

SEND ALL CORRESPONDENCE TO

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New PCT patent application  
Multhoff, Gabriele  
Our Ref.: D 1226 PCT/a

**English translation of the PCT application****New Use of Hsp70 Protein**

The invention relates to the use of Hsp70 protein or fragments thereof for the activation of NK-cells, pharmaceuticals, medical products or medical adjuvants which contain a Hsp70 protein or fragments thereof or activated NK-cells, a method for the activation of NK-cells as well as medical uses of the products obtained by the method of the invention.

Chaperones are necessary for a number of fundamental processes in the cell. In particular, it is known that they counteract cell stress. The best analysed class of chaperones is the group of the heat-shock proteins (HSP) with a molecular weight of 70 kDa (Multhoff et al., Cell Stress & Chaperones 1 (3) (1996), 167). These proteins are highly preserved in an evolutionary way. They bind to unfolded or incorrectly folded polypeptides in the cell, stabilize them and in that way inhibit their aggregation or make it possible for a transmembrane translocation to take place. Hsp70 is localized in the nucleus, in the cytosol and on the cell surface of certain tumour cells. Apart from their intracellular task as chaperones the members of the Hsp70 family seem to play a role as to the stimulation of the immune system, e.g. inflammatory processes under participation of pathogens, and as to the cellular anti-tumour immune response in vivo and in vitro. Accordingly, a range of therapeutic uses of heat-shock proteins has been presented in the state of the art. In WO 97/10000, the use of complexes consisting of a heat-shock protein and an exogenous antigen molecule (peptide) bound to this protein in a non-covalent way is described for the prevention and treatment of tumour and infectious diseases. The antigens present with the heat-shock proteins in the complex derive from tumour cells. They have the same characteristic, i.e. both induce an immune response. Multhoff et al., Biol. Chem. 379 (1998), 295-300 assign a role to HSPs, including Hsp70, in the

recognition of non-MHC-restricted effector cells, including NK-cells. In particular, it was found that NK-cells recognise Hsp70 molecules present on the surface of tumour cells and then lyse the tumour cells. Another approach of therapy for cancer diseases has been introduced by Tamura and colleagues (Tamura et al., Science 278 (1997), 117-123). They were able to demonstrate that tumour-carrying mice can be treated successfully with heat-shock preparations if derived from autologous tumours. Preparations from non-autologous tumours or from normal tissue, however, do not lead to the regression of the tumours (Blachere et al., J. Exp. Med. 186 (1997) 1315-1322). In the study by Tamura et al., the HSPs form complexes with a number of peptides which have not been identified in more detail. To sum up one can say that the state of the art shows an immunological activity of HSP molecules if they either form complexes with peptides and/or are present on the surface of cells such as tumour cells. Thus, although the potential of heat-shock proteins basically has been recognised for fighting various diseases, the successful therapeutic use usually depends on the preparation of certain complexes or cell preparations as well as the amount of the starting material (tumour material): It is hard to imagine a universal or patient-independent use of these complexes or cell preparations.

Thus, the problem underlying the present invention was to provide new means and methods for the use of the immunological potential of heat-shock proteins which do not have the above-mentioned disadvantages of the state of the art.

This problem has been solved by providing the embodiments characterised in the claims.

Therefore, the invention relates to the use of a Hsp70 protein, a carboxy-terminal (C-terminal) fragment thereof or a derivative thereof or a protein with an amino acid sequence homology to the C-terminal region of the Hsp70 protein of  $\geq 70\%$  for the production of a pharmaceutical preparation, medical product or a medical adjuvant for the activation of NK-cells.

According to the invention, pharmaceutical preparations are defined as substances and preparations of substances which, when used on or in the human body, are meant for healing, alleviating, preventing or recognising diseases, ailments, physical defects or pathological discomforts.

According to the invention, medical products are all substances or preparations used individually or in combination with each other of substances, or other subject-matters which, according to the producer, are meant to be applied to humans due to their functions for the purpose of detecting, preventing, monitoring, treating or alleviating

diseases and whose main effect in or on the human body is achieved neither by pharmacologically or immunologically effective preparations nor by a metabolism whose effectiveness may well be supported by such preparations.

According to the invention, medical adjuvants are such substances which are used for the production (as active ingredients) of pharmaceutical preparations.

Moreover, the invention relates to the use of a Hsp70 protein, a C-terminal fragment thereof or a derivative thereof or a protein with an amino acid sequence homology to the C-terminal region (amino acids 384-641) of the Hsp70 protein of  $\geq 70\%$  for ex vivo or in vitro activation of NK-cells.

According to the invention, the term "Hsp70 protein" relates to eukaryotic heat-shock proteins, the expression of which can be induced by heat but also by a number of other reagents such as e.g. amino acid analogues, heavy metals, ionophores or cytotoxines, wherein the factor of the increase in the expression by means of induction is at least 5, compared to the constitutive expression. The enclosed Figure 5 shows the composition of a Hsp70 protein comprising an N-terminal ATPase domain and a C-terminal substrate-binding domain of the protein. The complete amino acid sequence has been published in Milner et al., Immunogenetics 32 (4) (1990), 242-251.

According to the invention, the term "carboxy-terminal (C-terminal) fragment" of the Hsp70 protein comprises (poly)peptides exhibiting an amino acid sequence from the range of amino acids 384-641 of the human Hsp70. The present invention also comprises fragments of the C-terminal fragment 384-641. In further embodiments the term comprises (poly)peptides derived from the region of another protein that is also comprised in the term "Hsp70 protein" used according to the invention, this region being homologous to the above-mentioned C-terminal region of the human Hsp70 protein. The fragments of the Hsp70 protein used according to the invention are also able to activate NK-cells. The activation can easily be verified by the person skilled in the art by means of the teaching of the invention. Thus, the person skilled in the art is also able to produce fragments from the above-mentioned fragment 384-641 by recombinant techniques without further ado (standard methods for this are described in Sambrook et al., "Molecular Cloning, A Laboratory Manual", 2. edition 1989, CSH Press, Cold Spring Harbor, N. Y.) and test them for the activation properties wanted.

The term "derivate" comprises both derivatives of the Hsp70 protein as well as derivatives of the C-terminal fragment as far as the derivatives exhibit the functions of the invention. Preferably, such derivatives exhibit the same three-dimensional structure as Hsp70 or its

C-terminal fragment and can be produced, for example, by peptidomimetics (al-Obeidi et al., Mol. Biotechnol. 9 (1998), 205-223; Wiley et al., Med. Res. Rev. 13 (1993), 327-384; Bohm, J. Comput. Aided Mol. Des. 10 (1996), 265-272; Hruby et al., Biopolymers 43 (1997), 219-266).

The term "NK-cells" ("natural killer cells") comprises big, granular lymphocytes expressing CD45 on the surface and exhibiting killer activity without prior stimulation. They are particularly characterised in that they express CD16 and/or can be stimulated by interleukin-2 and/or do not express CD3 and/or do not have  $\alpha/\beta$ - or  $\gamma/\delta$ -T-cell receptors.

The NK-cells developing their effectiveness in the method of the invention are further characterised by the following properties:

- they are transient plastic-adherent after addition of IL-2 in amounts of 10 to 10,000 Units, e.g. of 100 I U, wherein IL-2 can be purchased from the firm Chiron;
- the adherence takes effect 3-18 hours after addition of IL-2 on newly isolated PBL (peripheral blood lymphocytes depleted by monocytes);
- the NK-cells exhibit a CD16<sup>dim</sup> expression (average value of fluorescence weak);
- the NK-cells express CD56 and CD57 as typical NK marker;
- the NK-cells express CD94 (C-type lectin killer cell receptor)
- the NK-cells secrete after activation with Hsp70 and cytokins IFN $\gamma$ ;
- the NK-cells can be stimulated by addition of Hsp70 (purified protein) (growth and cytotoxic activity);
- they are not dependent on the patient's MHC type.

According to the invention, other NK-cell populations can be used, too. In this case, however, it is a pre-requisite that they can be activated by the Hsp70 used according to the invention or by the above-mentioned fragments or derivatives. According to the invention, isolated NK-cells can be used. It is furthermore possible to use cell mixtures such as peripheral mononucleic blood cells (PBMC) containing NK-cells.

According to the invention, the term "amino acid sequence homology to the C-terminal region of the Hsp70 protein  $\geq 70$  %" means that at least 70 % of the amino acids are identical when two amino acid sequences are aligned, wherein one of the aligned amino acid sequences is the one of the C-terminal region of Hsp70. It also comprises sequences which have 70 % identical amino acids but which, in addition, however, differ from the C-terminal Hsp70 reference sequence by gaps when aligned. These gaps can occur either in the homologous molecule used according to the invention or in the reference molecule. Alignments, usually by comparison by computer, are known in the

state of the art, as are the programs with which such alignments can be carried out. Furthermore, it is preferred that the proteins or the fragments exhibit an amino acid sequence homology to the carboxy-terminal region, i.e. in the region of the amino acids 384 – 641, of the Hsp70 protein of  $\geq 80\%$  and preferably  $\geq 90\%$ .

The finding that heat-shock proteins, C-terminal fragments thereof or derivatives derived therefrom induce immunological activities by means of activation of NK-cells even if they do not form complexes with peptides or if they are not presented as tumour cells on the surface of cells has to be considered most surprising. In June 1998 (cf. Srivastava et al., Immunity 8 (1998), 657-665, herein cited as expert opinion), for example, it was still assumed that isolated heat-shock proteins do not exhibit immunogenic effects such as CTL induction (Blachere et al., J. Exp. Med. 186 (1997), 1315-1322) and cannot induce protective immunity against some kind of cancer (Udono and Srivastava, J. Immunol. 152 (1994), 5398-5403). With the present invention it becomes possible to induce an in vitro or in vivo activation of NK-cells which is independent from the patient and which can be used successfully against various diseases, for example, tumour diseases. Moreover, the use of isolated heat-shock protein allows for an improved standardisation of activation processes. The present invention makes it possible to produce unlimited amounts of HCP, whereas in the case of patient-specific preparations of the HSP peptide complexes the amount of HSP is limited by the size of the tumor.

In addition to the above-mentioned findings it is surprising that the use of C-terminal fragments of the Hsp70 protein also leads to the result of the invention. In particular, it was unexpected that the C-terminal apparently exhibits the same three-dimensional structure which is recognized by the NK-cells and which leads to their activation. The possibility of using the C-terminal Hsp70 fragments in the activation of NK-cells also has the advantage that the recombinant production should lead to increased yields in comparison to the recombinant presentation of the whole protein.

The production of derivatives of Hsp70 or its C-terminal fragments, e.g. by peptidomimetics, is for example useful if a rapid degradation of these (poly)peptides in the body is to be avoided. This can play a role e.g. with oral administration of pharmaceutical preparations.

Preferably, the activation comprises the induction of an immune response mediated by NK-cells. Here, it is particularly preferred that the immune response mediated by NK-cells comprises a stimulation of the proliferation of the NK-cells and/or an increase of the cytolytic activity of the NK-cells. (Pharmaceutically) effective amounts of Hsp70 or of fragments or derivatives thereof are used in all of the above-mentioned embodiments so

that the desired activation, preferably the immune response, is induced. The immune response is primarily, but not exclusively, directed against such cells expressing Hsp70 or fragments thereof on the cell surface. This includes both human and animal cells. Human or animal cells expressing Hsp70 on the cell surface include, for example, tumour cells and cells from patients with infectious diseases. The cytolytic activity of the NK-cells stimulated by Hsp70 according to the invention is significantly increased so that an immunological elimination of these cells expressing Hsp70 on the cell surface becomes possible.

In a particularly preferred embodiment of the use according to the invention the cytolytic activity against tumour cells and/or cells from patients with infectious diseases is increased.

In another particularly preferred embodiment of the use according to the invention the cytolytic activity against leukaemia cells, lymphoma cells, tumour cells and metastasizing cells of solid tumours and cells from patients with viral, mycotic or bacterial infectious diseases is increased. One example of the treatment of viral diseases is the treatment of HIV infections, an example of a bacterial infection is the treatment of diseases caused by mycobacteria. Solid tumours whose metastatic cells can be treated by the immunological method of the invention include, for example, carcinomas, sarcomas, melanomas, leukaemias and lymphomas. Examples of carcinomas are colon carcinomas and lung carcinomas.

By using a Hsp70 protein, parts of this protein or  $\geq 70\%$  sequence homologous proteins according to the invention, cells which have been infected by viruses, bacteria and/or fungi or cells which have been changed by tumours can be lysed. Cells containing antigenic parts of these foreign organisms or parts of tumour cells can also be lysed by means of activated NK-cells by the use of the Hsp70 protein according to the invention.

The invention further relates to a method for ex vivo or in vitro activation of NK-cells, wherein a physiological cell suspension containing NK-cells is mixed with a Hsp70 protein, a C-terminal fragment thereof or a derivative thereof or a protein with an amino acid sequence homology to the C-terminal region (amino acids 384-641) of the Hsp70 protein of  $\geq 70\%$  and incubated to bring about an activation of the NK-cells.

The incubation can take place at room temperature, preferably, however, at physiological temperature (37°C) on a shaker (gentle shaking).

In a preferred embodiment of the method according to the invention the activation comprises a stimulation of the proliferation of the NK-cells and/or an increase in their



cytotoxicity. With regard to the preferred target cells for the cytotoxic activity it is referred to the above-mentioned explanations.

In a particularly preferred embodiment of the method of the invention peripheral, mononucleic blood cells (PBMC) or a fraction thereof which contain NK-cells are used as physiological cell suspensions containing NK-cells.

Using appropriate methods, the NK-cells can be obtained from the patients to be treated or from a healthy donor by taking blood. Preferably, buffy-coats (lymphocyte concentrates) containing NK-cells are to be used.

Buffy-coats (lymphocyte concentrates) are taken from patients via the veins and e.g. heparin is added to prevent clotting of the cells. The buffy-coats to which heparin has been added are gathered in a sterile receptacle (usually little plastic bags) and then centrifuged resulting in an accumulation of blood cells (= PBMC, peripheral, mononucleic blood cells, e.g. lymphocytes, erythrocytes, granulocytes, and so on). The lymphocyte concentrate remains sterile in the vessel (plastic bag). In the case of healthy probands a buffy-coat consists of white and red blood cells (lymphocytes, erythrocytes, and so on). In the case of a tumour patient the buffy-coat does not only consist of blood cells but can also contain tumour cells (in the case of leukaemia, e.g. leukaemia cells = blasts; in the case of solid tumours, e.g. metastasized cells).

The buffy-coats containing peripheral, mononucleic blood cells are used in the form of a physiological cell suspension, preferably with heparin added. The heparin prevents an aggregation of the cells.

In another particularly preferred embodiment of the method of the invention the cell suspension further contains human and animal cells expressing Hsp70 on the cell surface. Stimulation of the NK-cells by the Hsp70 protein however, can also take place without target cells (tumour cells, infected cells) that express Hsp70 on the cell surface being present.

In another particularly preferred embodiment of the method tumour cells, cells from patients with infectious diseases are used as human or animal cells.

In another particularly preferred embodiment of the method leukaemia cells, lymphoma cells, metastasizing cells of solid tumours and cells from patients with viral, mycotic or bacterial infectious diseases are used as human or animal cells.

In another preferred embodiment of the method the physiological cell suspension containing the cells and proteins is incubated for at least 3 hours.

In this embodiment, in order to increase the cytolytic effect of the natural killer cells, the target cells of the natural killer cells are preferably incubated together with the natural killer cells and the Hsp70 in suspension, preferably for the aforementioned period. Long-term incubations, however, are also possible for at least 4 days. Thus, in another particularly preferred embodiment of the method according to the invention, incubation is carried out for 4 days.

In another preferred embodiment of the use according to the invention or of the method according to the invention a cytokine is used in addition. The cytokine and the NK-cells and/or the heat-shock proteins, fragments or derivatives thereof can be used separately or together in one dose.

In a particularly preferred embodiment of the use or the method an interleukin is used as cytokine. According to the invention, also a combination of interleukins can be used with the Hsp70 protein to further enhance the activation of the NK-cells, e.g. the immune response mediated by the NK-cells or the stimulation of the proliferation of the NK-cells.

In another particularly preferred embodiment of the use or the method according to the invention IL-2, IL-12 and/or IL-15 is used as interleukin.

The invention makes it possible to re-infuse the patient not only with ex-vivo-activated NK-cells, but also to use in vivo NK-cells treated according to the invention due to avoidance of toxic substances, e.g. in combination with a hyperthermia treatment. This embodiment of the invention has a further, invaluable and surprising advantage, i.e. that the target cells, too, e.g. tumour cells which were resistant to the known therapeutic methods, can now be killed immunologically by the cytolytic effect of the NK-cells. Thus, the invention further relates to a method for in-vivo activation of the immune system, wherein a pharmaceutically effective amount of NK-cells activated according to the above-mentioned method is administered to a patient, optionally in combination with or before a pharmaceutically effective amount of a Hsp70 protein, a C-terminal fragment thereof or a derivative thereof or a protein with an amino acid sequence homology to the C-terminal region (amino acids 384-641) of the Hsp70 protein of  $\geq 70\%$ .

If substances are administered together, they may be put together in one container or separately in several containers, whereas in the case of consecutive processing they are kept separately.

If the NK-cells are administered before the Hsp70 proteins, the corresponding time period before administration of the Hsp70 protein should be at least 3-24 hours.

An example of treatment according to the invention is the following:

Buffy-coat cells (lymphocyte concentrates) consisting of peripheral, mononucleic blood cells or bone marrow cells and tumour cells from tumour patients, for example leukaemia patients, are subjected to heat treatment in a sterile sealed container, for example a plastic container, with the Hsp70, Hsp70 related protein and/or effective fragments or derivatives thereof in a water bath with temperature control. Both the tumour cells and the NK-cells stimulated by the present treatment are in the container. After completion of the procedure of the invention, the culture solution containing the activated NK-cells and the lysed tumour cells is re-infused into the patient.

According to the invention the NK-cells are optionally present with other peripheral, mononucleic blood cells, for example with erythrocytes and granulocytes and T-cells. Preferably, therefore, the NK-cells are not used alone, but a mixture of the peripheral, mononucleic blood cells is obtained by isolating buffy-coat cells. These accumulations further contain tumour cells in tumour patients which are immunologically eliminated by the method of the invention.

Furthermore, the invention relates to a method for in vivo activation of NK-cells, wherein a patient is given a pharmaceutically effective amount of a Hsp70 protein, a C-terminal fragment thereof or a derivative thereof or a protein with an amino acid sequence homology to the C-terminal region (amino acids 384-641) of  $\geq 70\%$ .

Moreover, the invention relates to a method for the treatment of tumours, cancer diseases and/or infectious diseases, wherein a patient is given a pharmaceutically effective amount of NK-cells activated according to the above-mentioned method of the invention and/or a Hsp70 protein, a C-terminal fragment thereof or a derivative thereof or a protein with an amino acid sequence homology to the C-terminal region (amino acids 384-641) of the Hsp70 protein of  $\geq 70\%$ .

In a preferred embodiment of the method of the invention the tumour is a solid tumour or a metastasis. The treatment strategy particularly aims at the elimination of single-cell metastases which can be eliminated by the method of the invention. An increased activity of Hsp70-specific cells can be achieved by addition of interleukin-2 in a low dosage, for example 100 I.U.. The interleukin-2 may, for example, be introduced into the sterile container, e.g. a plastic container, with the Hsp70.

In another preferred embodiment of the invention the cancer disease is leukaemia or a lymphoma.

In another preferred embodiment of the invention the infectious disease has a viral, mycotic or bacterial origin.

The invention further relates to a pharmaceutical preparation, a medical adjuvant, or a medical product containing a Hsp70 protein, a C-terminal fragment thereof or a derivative thereof or a protein with an amino acid sequence homology to the C-terminal region (amino acids 384-641) of the Hsp70 protein of  $\geq 70\%$  and/or of NK-cells activated according to the method of the invention in a therapeutically effective amount as well as optionally common carrier and/or adjuvant substances. Furthermore, a cytokine as defined above is optionally added to the pharmaceutical preparation.

In a preferred embodiment of the pharmaceutical preparation, the medical adjuvant, or the medical product, the protein is present in a concentration of at least  $1\text{ }\mu\text{g/ml}$ , preferably up to  $1000\text{ }\mu\text{g/ml}$ , preferably  $1 \times 10^6$  to  $5 \times 10^8$  NK-cells, wherein an amount of  $10\text{ }\mu\text{g}$  to  $600\text{ }\mu\text{g/ml}$  is preferred.

Examples of suitable pharmaceutically tolerable carriers are known to the person skilled in the art and comprise, for example, phosphate-buffered saline solutions, water, emulsions, such as oil/water emulsions, sterile solutions, and so on. The pharmaceutical compositions (pharmaceutical preparations) containing such carriers may be prepared according to common methods. The pharmaceutical compositions may be administered to the respective individuals in an appropriate dosage. Ways of administration are, for example, intravenous, intraperitoneal, subcutaneous, intramuscular, topic or intradermal. The dosage depends on many factors, e.g. on the patient's size, sex, weight, age as well as the type of the composition specially administered, the kind of administration and so on. In general, the dosage given per month is  $10$  to  $1000\text{ }\mu\text{g}$ . In connection with the intravenous injection of substances of the invention dosages of  $10$  to  $1000\text{ }\mu\text{g}$  are usual. The compositions may be administered locally or systemically. Generally, administration is carried out parenterally. Therefore, the NK-cells treated with Hsp70 according to the invention are preferably injected intravenously. An injection may also be carried out directly into the tumour with an effective amount of NK-cells being injected. Other known types of application are, of course, also possible.

The Hsp70 itself may, for example, be applied with cytokines. One example of application is the injection of the Hsp70 protein, e.g. with cytokines in an intravenous, intramuscular, a subcutaneous or an intraperitoneal way or, also in the sole of the foot.

In another preferred embodiment of the use or the method or the pharmaceutical preparation or the medical product or the medical adjuvant of the invention, the Hsp70 protein is a human protein. The protein of the invention is, for example, of human origin (with the isolation of cell extracts) or exhibits the amino acid sequence of the human Hsp70 protein (e.g. after recombinant production). Animal Hsp70 proteins may, however, also be used.

The Hsp70 protein or the fragments used according to the invention may be produced recombinantly, isolated from cell extracts, or by means of chemical synthesis.

Preferably, the Hsp70 protein or its fragment or derivative is a recombinant protein. Such recombinant proteins can be produced according to standard techniques.

Such standard techniques are known to the person skilled in the art (Sambrook et al., *ibid.* and Ausubel, "Current Protocols in Molecular Biology", Green Publishing Associates and Wiley Interscience, N.Y. (1989)). For the recombinant production of proteins nucleic acid molecules are used which encode the Hsp70 protein or fragments thereof. These may be various nucleic acid molecules, in particular DNA or RNA molecules, for example cDNA, genomic DNA, mRNA and so on. Such nucleic acid molecules may be naturally occurring molecules and/or molecules produced by genetic or chemical synthesis methods. For the production of recombinant proteins, the person skilled in the art uses vectors, particularly plasmids, cosmids, viruses, bacteriophages and other vectors common in genetic engineering (Sambrook et al., *ibid.*). The nucleic acid molecules contained in the vectors may be linked to regulatory elements which guarantee the expression in prokaryotic or eukaryotic cells. In this context, the term "expression" can mean both transcription and transcription and translation. Regulatory elements comprise promoters, in particular. For the expression of a nucleic acid molecule in prokaryotic cells a range of promoters are at disposal, e.g. the *E. coli* lac- or trp-promoter, the  $P_R$ - or  $P_L$ -promoter of the lambda-phage, lacI, lacZ, T3, T7, gpt, and so on. Eukaryotic promoters are, for example, the CMV immediate early promoter, the HSV promoter, the thymidinekinase promoter, the SV40 promoter, LTRs from retroviruses and the mouse metallothionin I-promoter. A variety of expression vectors for the expression in prokaryotic or eukaryotic cells has been described, e.g. for eukaryotes pKK223-3 (Pharmacia Fine Chemicals, Uppsala, Sweden) or GEM1 (Promega Biotec, Madison, WI, USA), pSV2CAT, pOG44 and for prokaryotes pQE70, pQE60, pBluescript SK, and so on. Apart from promoters these vectors can also contain elements for further enhancement of the transcription, such as the so-called transcription enhancers. Examples are the SV40 enhancer, the polyoma enhancer, the cytomegalovirus early promoter enhancer and adenovirus enhancer. The recombinant proteins may, therefore, be expressed in various

prokaryotic or eukaryotic host cells, for example, when the above-described vectors are used. Examples of such host cells are bacterial cells (such as the E. coli, streptomyces, bacillus, salmonella typhimurium), fungus cells (such as yeast cells, particularly saccharomyces cerevisiae), insect cells (such as drosophila or SF9 cells), animal cells (such as CHO or COS cells) or, also, plant cells and so on. Such host cells are cultivated under conditions allowing for the expression of the recombinant protein which may then be gained from the cells and/or from the culture medium. Methods for the expression of foreign protein in various types of host cells as well as for gaining the produced protein are known to the person skilled in the art.

In another preferred embodiment of the use or the method or the pharmaceutical preparation or the medical product or the medical adjuvant of the invention, the Hsp70 protein comprises the C-terminal fragment (amino acids 384 to 561) of the human Hsp70 or the corresponding region of another Hsp70 exhibiting the effects of the invention, or the C-terminal fragment (amino acids 454 to 460) of the human Hsp70. According to the invention it could surprisingly be shown that fragments having this minimal sequence of 7 amino acids (NLLGRFE) are inhibited by antibodies, and NK activation was thus prevented or stopped. The experiments were carried out according to Multhoff et al., J. Immunol. 158 (1997), 4341-4350. The antibody RPN 1197 available from Amersham was used. The 7 amino acids can be flanked by naturally flanking Hsp70 sequences or by other amino acids. The 7 amino acids are preferred to remain in their natural context. When other flanking amino acids are used, the three-dimensional context, in which said 7 amino acids naturally occur, is preferred to be maintained. Within the 7 amino acids further amino acid exchanges can take place as far as the homology of at least 70 % is maintained. These exchanges, however, do not comprise an exchange of arginine in position 458 by lysine. This exchange leads to a change in confirmation. Accordingly, exchanges in the region of the 7 amino acids which lead to a change of confirmation are comprised by the invention only if they exhibit the activation properties desired.

The invention further relates to the use of the NK-cells treated according to a method of one or more of the above-mentioned embodiments for therapy of tumour diseases and/or infectious diseases.

In a preferred embodiment of the use of the invention the therapy is carried out by re-infusion of the NK-cells treated.

The aforementioned in vivo methods may also be used as treatment methods for treating the described indicators.

The Figures show:

**FIGURE 1:** Comparison of the proliferating activity of separated NK- (A)- and T (B) – cells which were stimulated at a concentration of 10 µg/ml each either with IL-2(100 IU/ml)–medium or with other recombinant Hsp70 proteins (rHsp70, rHsp70-C<sub>term</sub> (amino acids 384-561), rHsp70homC, DnaK, Hsc70 and heat denatured rHsp70) which are suspended in IL-2-medium (100 IU/ml). The phenotypic characteristics of the NK-cells are the following:

CD3: < 5 %; CD16/CD56: 46-87 %; CD94: 60-70 %; p58.1 and p58.2: < 5 % and T-cells: CD3: 85-92 %; CD16/CD56: 5-10 %; CD94: < 29 %; p58.1 and p58.2: not tested; p70: not tested, determined by flow cytometry. The proliferation of the cells was determined after 48 hours and after incubation with <sup>3</sup>H-thymidine (1 µCi/ml) at 37 °C for 18 hours. The relative percentage of the <sup>3</sup>H-thymidine absorption in NK – (A) and T (B) – cells was compared to the effects of IL-2 alone (100 %). The values show average values of four to seven independent experiments ± standard deviation.

**FIGURE 2:** Comparison of the cytotoxic activity of highly purified NK-cells (CD3: < 2 %; CD16/CD56; 75-80 %; CD94: 65-87 %; p58.1 and p58.2: 20-30 %; p70: < 10 %) which either remained untreated (continued lines, empty symbols) or which were pre-incubated after a pre-incubation of the NK-cells with rHsp70 (A) – protein (each 5 µg/ml for 4 days; continued lines, filled in symbols) compared to <sup>51</sup>Cr-labelled tumour target cells CX+ (A) and CX- (B) which are different due to their capability of express plasma membrane. The results are noted as percentage of the specific lysis at various E : T-ratios of 0.2 : 1 to 2 : 1. Each point represents the average value of at least three independent experiments ± standard deviation. The percentage of spontaneous release for each tumour target cell line was always below 15 %.

**FIGURE 3:** Tumour growth of CX+ cells carrying Hsp70 in immunodeficient mice. After i.p.-injection of NK-cells the tumour growth is completely inhibited (with i.p.-injection of the tumour cells). The tumour growth was given in cm<sup>2</sup>. Tumour growth after i.p.-injection of CX+ and NK-cells on day 21.

**FIGURE 4:** Tumour growth of CX+ cells carrying Hsp70 in immunodeficient mice. After i.v.-injection of NK-cells the tumour growth is completely inhibited. The tumour growth was measured in gram.

Tumor growth after o.t.-injection of CX+ and i.v.-injection of NK-cells on day 35. The NK-cells inhibit the tumour growth of CX+ cells carrying Hsp70 after 3 or 5 weeks after injection and (see Figure 3).<sup>1</sup> Both an intraperitoneal and an intravenous application of the NK-cells leads to similar results.

**FIGURE 5:** Intact Hsp70-protein with illustration of the C-terminal region

**FIGURE 6:** Illustration of the influence of Hsp70 and/or cytokines on the immune response mediated by NK-cells. The addition of cytokine also leads to a stimulation of T-cells.

Each of the documents cited herein (including instructions of the producers, manuals, etc.) has been incorporated by reference in the description.

The Examples illustrate the invention.

#### **Example 1: Increased proliferation of NK-cells after addition of Hsp70**

The proliferation of purified NK- and T-cells which were stimulated with the Hsp70 proteins rHsp70, DnaK, Hsc70, rHsp70-C term. and rHsp70homC (amino acids 384-561) was determined in a <sup>3</sup>H-thymidine absorption standard test (for test conditions see later). Additionally, first, peripheral blood lymphocytes from voluntary human donors were separated in non-adherent CD3+ T-cell and transient (12-24 hours) adherent CD3- (CD16+/CD56+) NK-cell-subpopulations in a multiple-steps method and subsequent 12-hour incubation in an IL-2-containing medium (see 3). The cells were cultivated separately in rIL-2 (100 IU, Chiron, Frankfurt, Germany) containing an RPMI 1640 (Life Technologies, Eggenstein, Germany) medium for 3-4 days. The proliferation was measured as H-3 absorption.

Panning experiments were carried out using human recombinant Hsp70 (rHsp70, SPP-755, StressGen Biotechnologies, Victoria, Canada) and DnaK (Hsp70-homologue obtained from E. coli, SPP-630, StressGen). The proteins were diluted in PBS to a stem concentration of 1 µg/ml and deep frozen in aliquots at -80°C. T-25 culture flasks were incubated with rHsp70 or DnaK proteins (10 µg/ml) diluted in 3 ml icy cold carbonate

<sup>1</sup> Translator's note: sentence incomplete.



buffer, pH 9.5, for 12 hours. After blocking non-specific binding sites with PBS/5 % FKS a mixture of T- and NK-cells suspended in PBS/1 % FKS at a ratio of 1 : 2 and 2 : 1 was incubated in the culture flasks for 1 hour at room temperature. Non-adherent cells were obtained from the supernatant fraction after incubation. Adherent cells were obtained by means of sequential washing steps using ice-cold PBS/10 % FKS solution. A single, mild washing step was applied to remove lightly-adherent cells, whereas highly-adherent cells were obtained by means of additional stringent washing steps. The cell populations obtained with each step were counted separately and characterised by flow cytometry and phenotype.

The flow cytometry was carried out on an FACScan instrument (Becton Dickinson, Heidelberg, Germany) as described in (4). The percentage of positive-dyed cells was defined as the difference between the number of specifically-dyed, vital (propidiumiodide-negative) cells minus the number of cells which were dyed with the control antibodies corresponding to the isotype. The following antibodies were used for the phenotypic characterisation:

The control antibody corresponding to the isotype (Dianova, Hamburg, Germany; Becton Dickinson, Heidelberg, Germany), CD16 (Dianova, Hamburg, Germany), CD3 (Dianova, Hamburg, Germany).

The ability of T- or NK-cells to proliferate against various Hsp70 proteins was determined in a  $^3\text{H}$ -thymidine absorption standard test. Viable cells ( $5 \times 10^4$  cells/100  $\mu\text{l}$ ) were put on a micro titer plate containing 96 wells with a flat bottom (Greiner, Nürtingen, Germany), wherein a supplemented RPMI 1640-medium with 100 IU IL-2 and various recombinant Hsp70 proteins (rHsp70, DnaK, Hsc70, the constitutive form of Hsp70, purified from bovine brain, SPP-750; Stressgen; rHsp70-C term. sie rek. C-terminal peptide binding domain of Hsp70 (amino acids 384-561), rHsp70homC, the recombinant C-terminal peptide binding domain of Hsp70hom (Hsp70hom is a testis-specific member of the Hsp70 family exhibiting a high homology (94 %) to Hsp70, amino acids 384-561) were used. By testing various concentrations of the Hsp 70 proteins (1 – 200  $\mu\text{g/ml}$ ) it was found that a final concentration of 100  $\mu\text{g/ml}$  was ideal for stimulation. A proliferating activity against IL-2 (100 IU) was determined in parallel as a further control. After an incubation period of 24 hours or 48 hours the cells were labelled with  $^3\text{H}$ -thymidine (1  $\mu\text{Ci/well}$ ) and the total absorption was determined after 18-hour incubation at 37°C in a fluid scintillation counter (Beckmann Instruments, Munich, Germany). Furthermore, as internal control, the proliferation capacity was determined from the T-lymphocytes taken from the same donor. Using various Hsp70 proteins/fragments in concentrations of 1-200  $\mu\text{g/ml}$  in dose escalation examinations showed that a maximum stimulation of the proliferation capacity could be achieved with 100  $\mu\text{g/ml}$  Hsp70 protein. The proliferation

activity of isolated NK- and T-cells was tested after in vitro stimulation with rHsp70, DnaK, Hsc70, rHsp70-C term. or rHsp70homC (amino acids 384-562). As can be seen in Figure 1A, the NK-cell proliferation by rHsp70 was significantly stimulated. The stimulation by the carboxy-terminal region of Hsp and by rHsp70homC which, in the C-terminal domain with the amino acids 384-561, is 94 % identical to Hsp70 is also possible. In contrast, DnaK and Hsc70 did not stimulate the proliferation of NK-cells. Heat denatured rHsp70 lost its stimulatory properties for the proliferation completely.

Moreover, it was possible to show that the proliferation of CD3-positive T-lymphocytes could be stimulated by DnaK whereas rHsp70, Hsc70 and rHsp70homC and heat denatured rHsp70 did not show any effect on the proliferation capability of T-cells (Figure 1B).

Thus, to sum up, a proliferation of NK-cells by recombinant human Hsp70 protein by means of the C-terminal region of Hsp70 (384-561) and rHsp70homC, a protein homologous to the Hsp70, could be induced while a proliferation of the T-cells could selectively be stimulated by bacterial Hsp70 (*E. coli* DnaK).

#### **Example 2: Increase of the cytolytic activity of NK-cells after addition of Hsp70**

A functional analysis of the NK-cells using Hsp70-expressing (CX+) and Hsp70-non-expressing (CX-) tumour cells showed that the plasma membrane expression by Hsp70 correlated with an increased sensitivity for the lysis mediated by NK-cells. This lysis mediated by NK-cells can be blocked by pre-incubation of the tumour cell lines with monoclonal antibodies directed against the carboxy-terminal region (amino acids 504-617) of Hsp70 and with the antibody RPN1197 (1, 4). In the following the influence of recombinant Hsp70 protein (rHsp70) on the cytolytic activity of NK-cells for the autologous, Hsp70-expressing (CX+) and Hsp70-non-expressing (CX-) tumour cells was analysed. The results of the cytotoxicity tests using highly-purified NK-cells pre-incubated with Hsp70 protein in a concentration of 5  $\mu\text{g/ml}$  for 4 days are summarised in Figures 2A and B. The experimental set-up was the following: For stimulation of the cytotoxic activity NK-cells were incubated with 10  $\mu\text{g/ml}$  rHsp70. The stimulation was repeated after 4 days.

The human, autologous colon carcinoma subcell lines CX+ and CX-, which differ in their Hsp70 expression on the plasma membrane (Multhoff et al., *J. Immunol.* 158 (1997), 4341), were cultivated in RPMI 1640 medium supplemented with 10 % FKS (Life Technologies), 6 mM L-glutamine and antibiotics (100 IU/ml) penicillin and 100  $\mu\text{g/ml}$

streptomycin (Life Technologies). Exponentially growing tumour cells were used as target cells and purified CD3- NK-cells were used as effector cells after cell sorting using a FACStar<sup>plus</sup> instrument (Becton Dickinson, Heidelberg, Germany). The cytotoxicity mediated by NK-cells was determined by carrying out a 4-hour <sup>51</sup>Cr-radioisotope test (Multhoff et al., J. Immunol. 158 (1997), 4341). The percentage of the specific lysis was calculated as follows: [(experimental release – spontaneous release) – maximum release – spontaneous release] x 100. The spontaneous release of Cr-51 was below 15 % in all the experiments.

Surprisingly, it was possible to demonstrate that both the proliferation and the cytolytic activity of NK-cells against Hsp70-expressing tumour cells (CX+) was stimulated when the NK-cells were incubated with rHsp70 protein for at least 4 days. In contrast, NK-cells from the same donor, which were not treated with rHsp70, lost this reactivity after 10 days (dates not shown). The lytic activity of NK-cells not stimulated with rHsp70 was lower compared to NK-cells treated and stimulated with rHsp70 and no significant difference in the lysis could be noted in the lysis of Hsp70-expressing and non-expressing tumour cells.

The present examinations show that the carboxy-terminal part of Hsp70 (amino acids 384-641) is responsible for the stimulation of cytolytic and proliferate function of NK-cells. According to the invention, it was thus possible to demonstrate that particularly the carboxy-terminal part of the Hsp70 protein is effective as stimulating signal for NK-cells which in vitro attack tumour cells specifically expressing the Hsp70.

### **Example 3: Anti-tumour effect with NK-cells stimulated by Hsp70**

For examining the in vivo relevance of NK-cells compared to Hsp70-expressing tumour cells tests were carried out on immunodeficient SCID/beige mice. First, various amounts of tumour cells (CX+ or CX- cells) were injected into SCID/beige mice. An amount of 2.5 mio. cells proved to be the optimal amount of tumour cells to induce tumour growth within a period of 3 to 5 weeks. An i.p. (intraperitoneal) or o.t. (orthotope, i.e. here, in the bowel wall) injection of colon carcinoma cells CX+ and CX- was chosen. The NK-cells were applied after stimulation either i.p. or i.v. (intravenously). As illustrated in Figure 3, tumour growth could be achieved in all animals both after i.p. and after o.t. injection. Contrary to the i.p. injection, in addition of growth of a primary tumour after o.t. injection metastasis of the CX+ could also be observed, in particular in spleen and lung (3 in 3 mice after o.t. injection).

Injection of human NK-cells (i.p., but also i.v.), even four days after the injection of tumour cells, leads to a complete inhibition of tumour growth. It is interesting that not only growth of primary tumour cells (in the i.p. region or in the bowels) could be inhibited by NK-cells but also the metastasis of tumours.

These findings clearly show that an immune reconstitution of SCID/beige mice with pre-activated human NK-cells does lead to lysis not only in vitro but also in vivo (in the animal). It is interesting that metastasis by human NK-cells, too, can be suppressed.

**English translation of the amended claims**

1. Use of a Hsp70 protein which does not form a complex with peptides when isolated together with these from tumour cells, a C-terminal fragment thereof or a derivative thereof or a protein with an amino acid sequence homology to the region of amino acids 384-641 of the Hsp70 protein of  $\geq 70\%$  for the production of a pharmaceutical preparation, a medical product or a medical adjuvant for the activation of NK-cells.
2. Use of a Hsp70 protein which does not form a complex with peptides when isolated together with these from tumour cells, a C-terminal fragment thereof or a derivative thereof or a protein with an amino acid sequence homology to the region of amino acids 384-641 of the Hsp70 protein of  $\geq 70\%$  for the in vitro or ex vivo activation of NK-cells.
3. Use according to claim 1 or 2, wherein the activation comprises the induction of an immune response mediated by NK-cells.
4. Use according to any one of claims 1 to 3, wherein the activation includes a stimulation of the proliferation of NK-cells and/or an increase of the cytolytic activity of NK-cells.
5. Use according to claim 4, wherein the cytolytic activity against tumour cells, cells from patients with infectious diseases is increased.
6. Use according to claim 5, wherein the cytolytic activity against leukaemia cells, lymphoma cells, tumour cells, metastasizing cells of solid tumours and cells of patients with viral, mycotic and/or bacterial infectious diseases is increased.
7. A method for the ex vivo or in vitro activation of NK-cells, wherein a physiological cell suspension containing NK-cells is mixed and incubated with a Hsp70 protein which does not form a complex with peptides when isolated together with these from tumour cells, a C-terminal fragment thereof or a derivative thereof or a protein

with an amino acid sequence homology to the region of amino acids 384-641 of the Hsp70 protein of  $\geq 70\%$  to effect activation of the NK-cells.

8. The method according to claim 7, wherein the activation comprises a stimulation of the proliferation of the NK-cells and/or an increase of their cytotoxicity.
9. The method according to claim 7 or 8, wherein peripheral, mononucleic blood cells or a fraction containing NK-cells is used as physiological cell suspension containing NK-cells.
10. The method according to any one of claims 7 to 9, wherein the cell suspension further contains human or animal cells expressing Hsp70 on the cell surface.
11. The method according to claim 10, wherein tumour cells, cells of patients with infectious diseases are used as human or animal cells.
12. The method according to claim 11, wherein leukaemia cells, lymphoma cells, tumour cells, metastasizing cells of solid tumours and cells of patients with viral, mycotic and/or bacterial infectious diseases are used as human or animal cells.
13. The method according to any one of claims 7 to 12, wherein the physiological cell suspension containing the cells and proteins is incubated for at least 3 hours.
14. The method according to claim 13, wherein the incubation is carried out for 4 days.
15. The method according to any one of claims 1 to 6 or the method according to any one of claims 7 to 14, wherein a cytokine is used in addition.
16. Use or method according to claim 15, wherein an interleukin is used as cytokine.
17. Use or method according to claim 16, wherein IL-2, IL-12 and/or IL-15 is used as interleukin.
18. A method for the in vivo activation of the immune system, wherein a patient is given a pharmaceutically effective amount of NK-cells activated according to the method according to any one of claims 7 to 17, optionally in combination with or before a pharmaceutically effective amount of Hsp70 protein, a C-terminal fragment thereof or a derivative thereof or a protein with an amino acid sequence homology to the region of amino acids 384-641 of the Hsp70 protein of  $\geq 70\%$ .

19. The method for the in vivo activation of NK-cells, wherein a patient is given a pharmaceutically effective amount of a Hsp70 protein which does not form a complex with peptides when isolated together with these from tumour cells, a C-terminal fragment thereof or a derivative thereof or a protein with an amino acid sequence homology to the region of amino acids 384-641 of the Hsp70 protein of  $\geq 70\%$ .
20. The method for the treatment of tumours, cancer diseases, infectious diseases or autoimmune diseases, wherein a patient is given a pharmaceutically effective amount of NK-cells activated according to the method according to any one of claims 7 to 17 and/or a Hsp70 protein which does not form a complex with peptides when isolated together with these from tumour cells, a C-terminal fragment thereof or a derivative thereof or a protein with an amino acid sequence homology to the region of amino acids 384-641 of the Hsp70 protein of  $\geq 70\%$ .
21. The method according to claim 20, wherein the tumour is a solid tumour or a metastasis.
22. The method according to claim 20, wherein the cancer disease is leukaemia or a lymphoma.
23. The method according to claim 20, wherein the infectious disease has a viral, mycological or bacterial origin.
24. Pharmaceutical preparation, medical product or medical adjuvant containing a Hsp70 protein, a C-terminal fragment thereof or a derivative thereof or a protein with an amino acid sequence homology to the region of amino acids 384-641 of the Hsp70 protein of  $\geq 70\%$  and/or NK-cells activated according to the method according to any one of claims 7 to 17 in a therapeutically effective amount as well as optionally common carrier substances and/or adjuvants except for Hsp70 protein which forms a complex with peptides when isolated together with these from tumour cells.
25. Pharmaceutical preparation, medical product or medical adjuvant according to claim 24, wherein the protein is present in a concentration of at least  $1\text{ }\mu\text{g/ml}$ , preferably up to  $1000\text{ }\mu\text{g/ml}$ .

26. Use according to any one of claims 1 to 6 or the method according to any one of claims 7 to 23 or pharmaceutical preparation, medical product or medical adjuvant according to claim 24 or 25, wherein the Hsp70 protein is a human protein.
27. Use according to any one of claims 1 to 6 or 26 or the method according to any one of claims 7 to 23 or 26 or pharmaceutical preparation, medical product or medical adjuvant according to any one of claims 24 to 26, wherein the Hsp70 protein or its fragment or derivative is a recombinant protein.
28. Use according to any one of claims 1 to 6, 26 or 27 or the method according to any one of claims 7 to 23, 26 or 27 or pharmaceutical preparation, medical product or medical adjuvant according to claim 24 to 27, wherein the Hsp70 protein comprises the C-terminal fragment (amino acids 384 to 561) of the human Hsp70 or the corresponding region of another Hsp70 comprising the effects of the invention.
29. Use of the NK-cells treated according to a method according to one or more of the above claims for the therapy of tumour diseases and/or infectious diseases.
30. Use according to claim 29, wherein the therapy is carried out by re-infusion of the treated NK-cells.

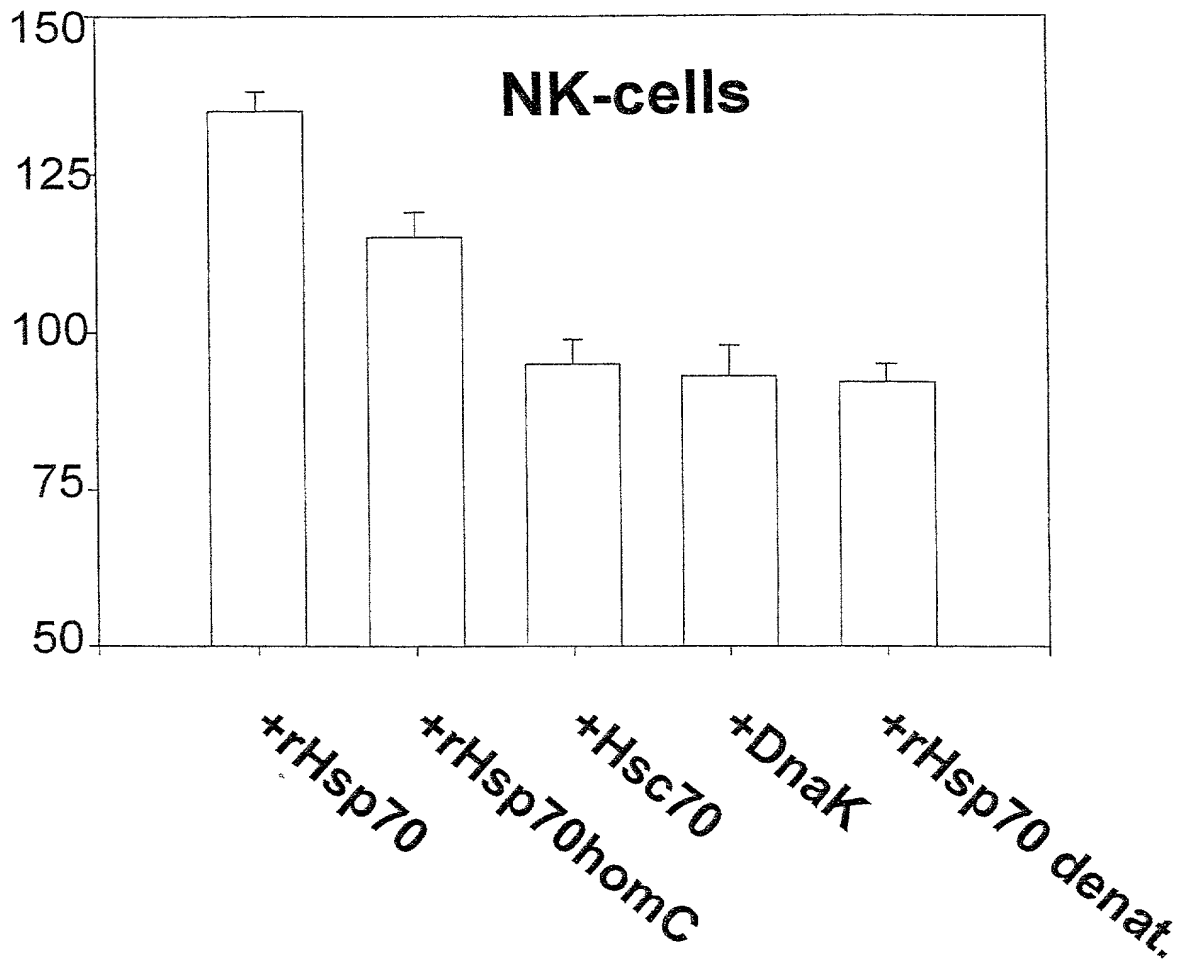


## **Abstract**

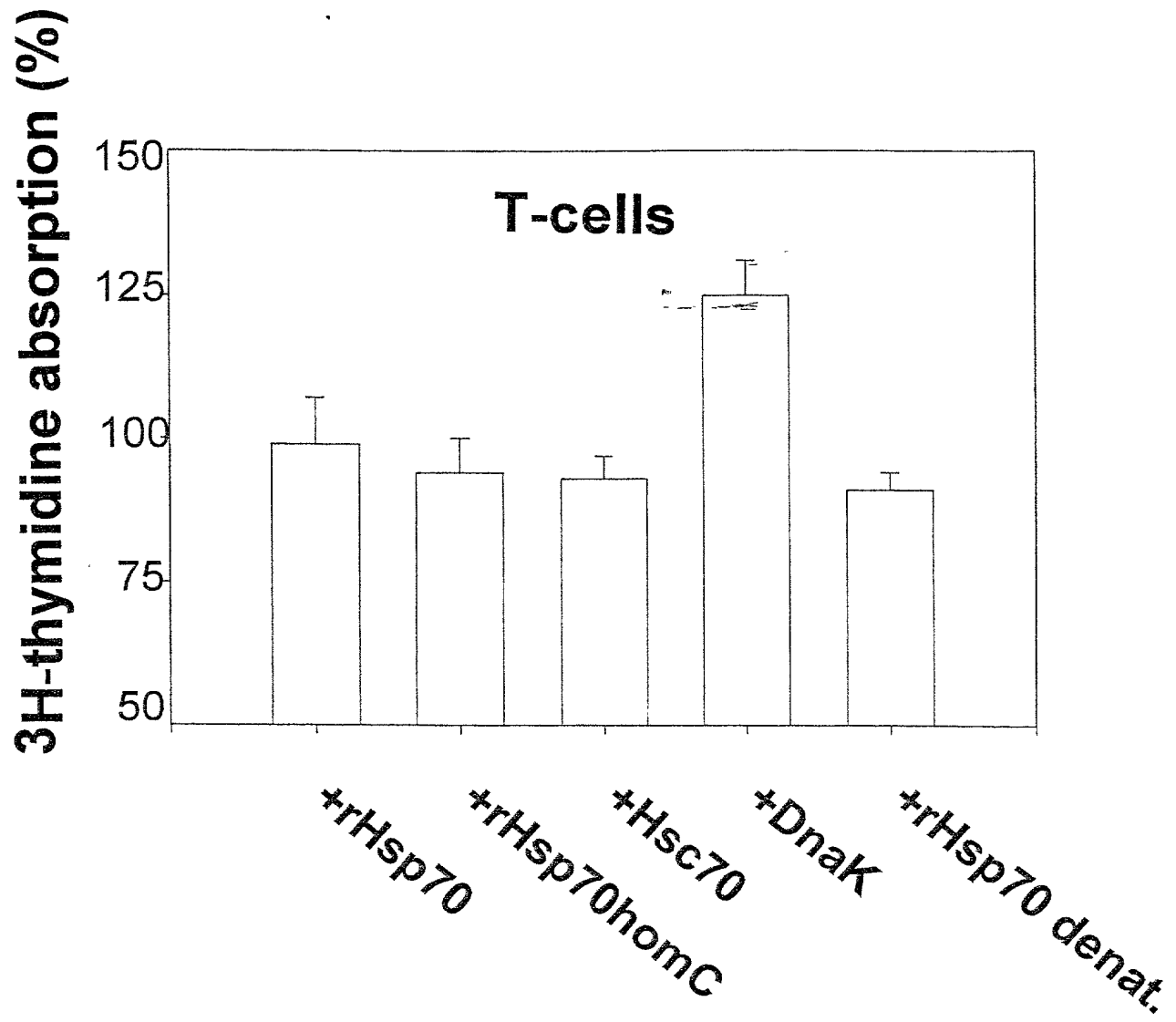
The invention relates to the use of Hsp70 protein or fragments thereof for the activation of NK-cells, pharmaceutical preparations, medical product or medical adjuvants containing a Hsp70 protein or fragments thereof or activated cells, method for the activation of NK-cells as well as medical use of the products obtained by the method of the invention.

FIG. 1A

**<sup>3</sup>H-thymidine absorption (%)**

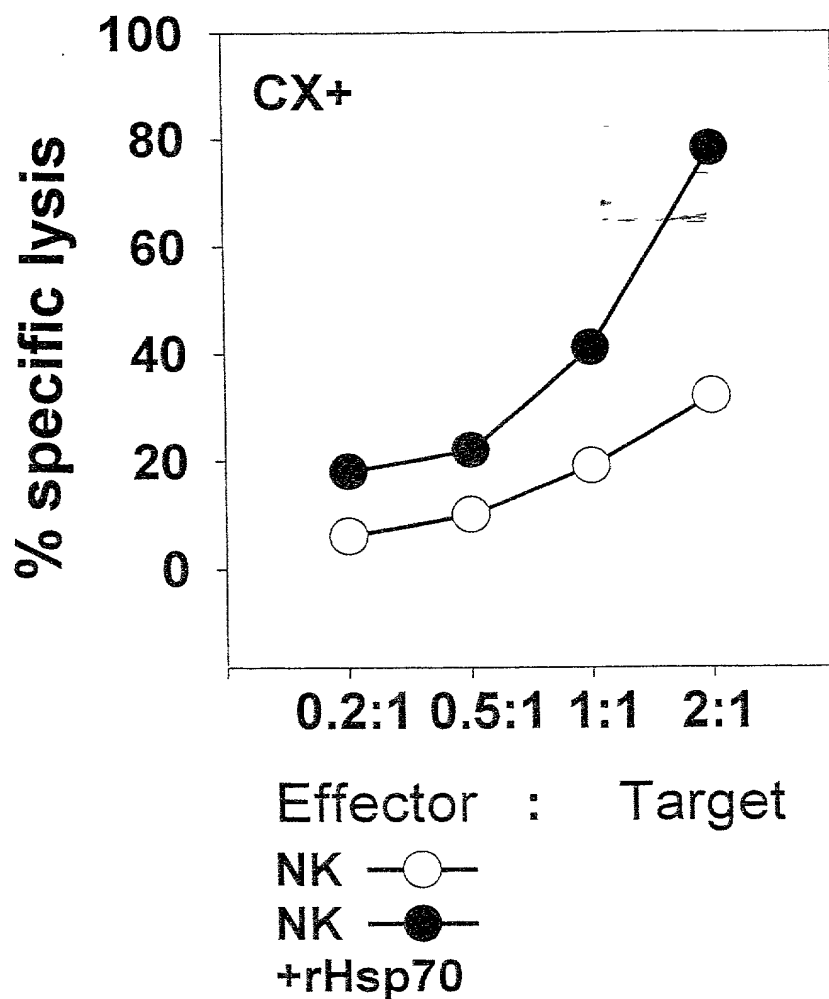


Growth of NK-cells increased by rHsp70, rHsp70-C<sub>term</sub> and rHsp70homC

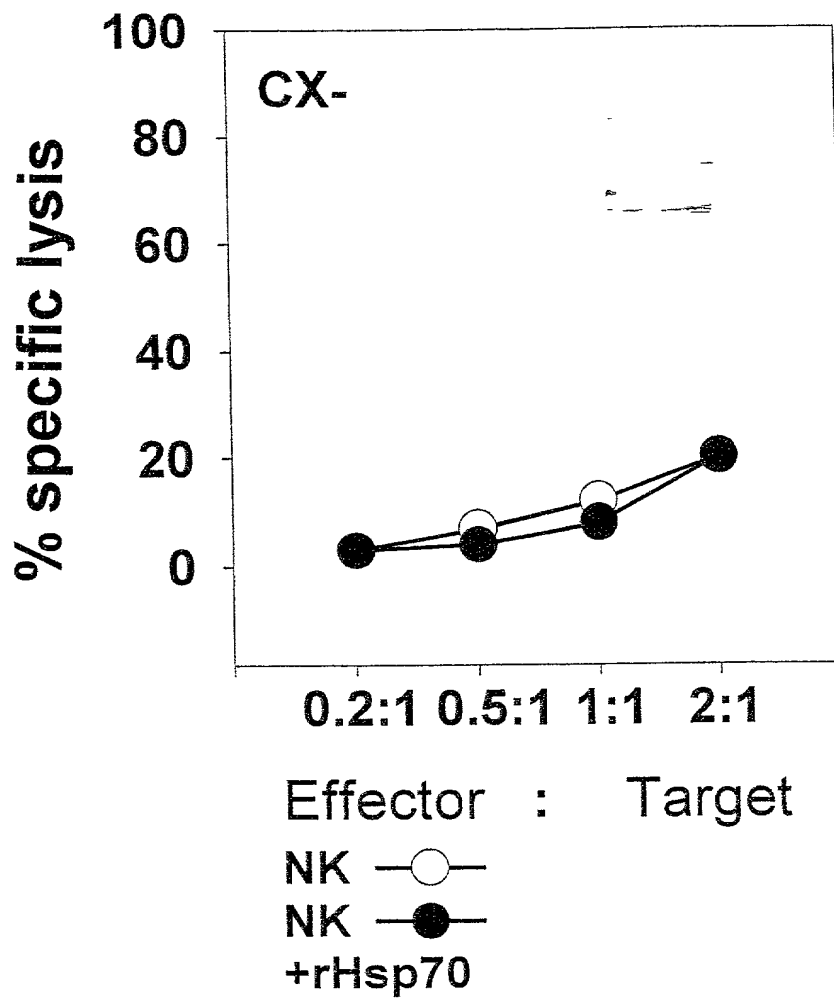
2/8  
FIG. 1B

Increase in the growth of T-cells exclusively by DnaK (=E. coli Hsp70)

FIG. 2A 3/8



Recombinant Hsp70-protein increases the lysis of CX+ tumour cells  
(which have Hsp70 on the membrane)

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FIG. 2B

Recombinant Hsp70-protein does not increase the lysis of CX- tumour cells (no Hsp70-membrane expression)

FIG. 3 5/8

Tumour growth after i.p. injection of CX+ and NK cells (day 21)

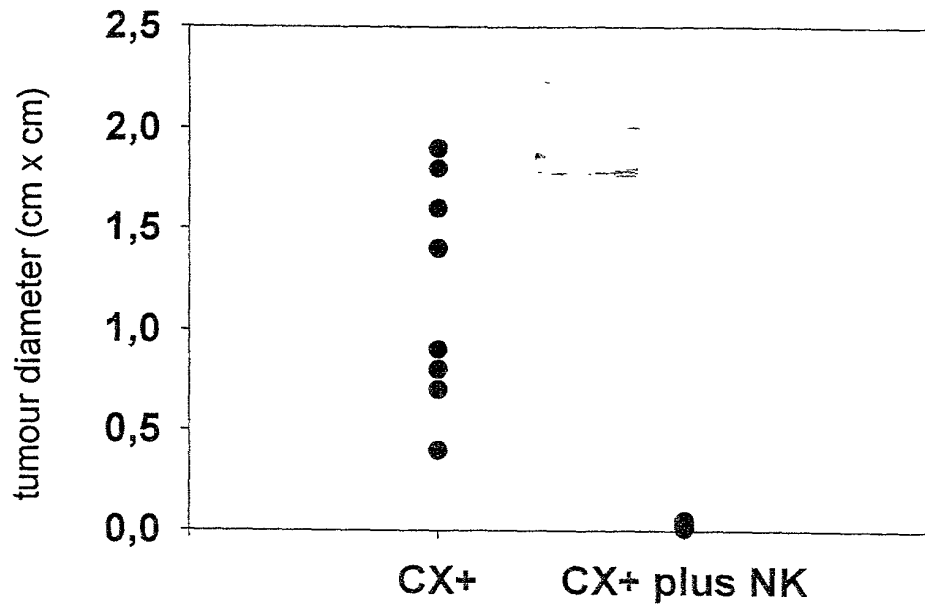


FIG. 4 6/8

Tumour growth after o.t. injection of CX+ and i.v. injection of NK cells (d35)

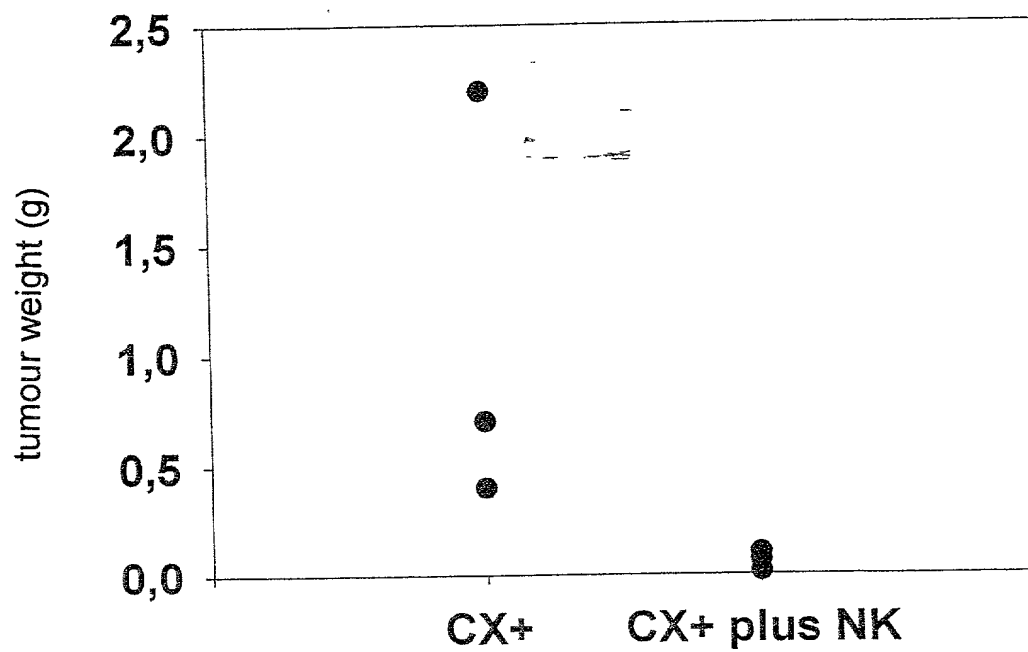
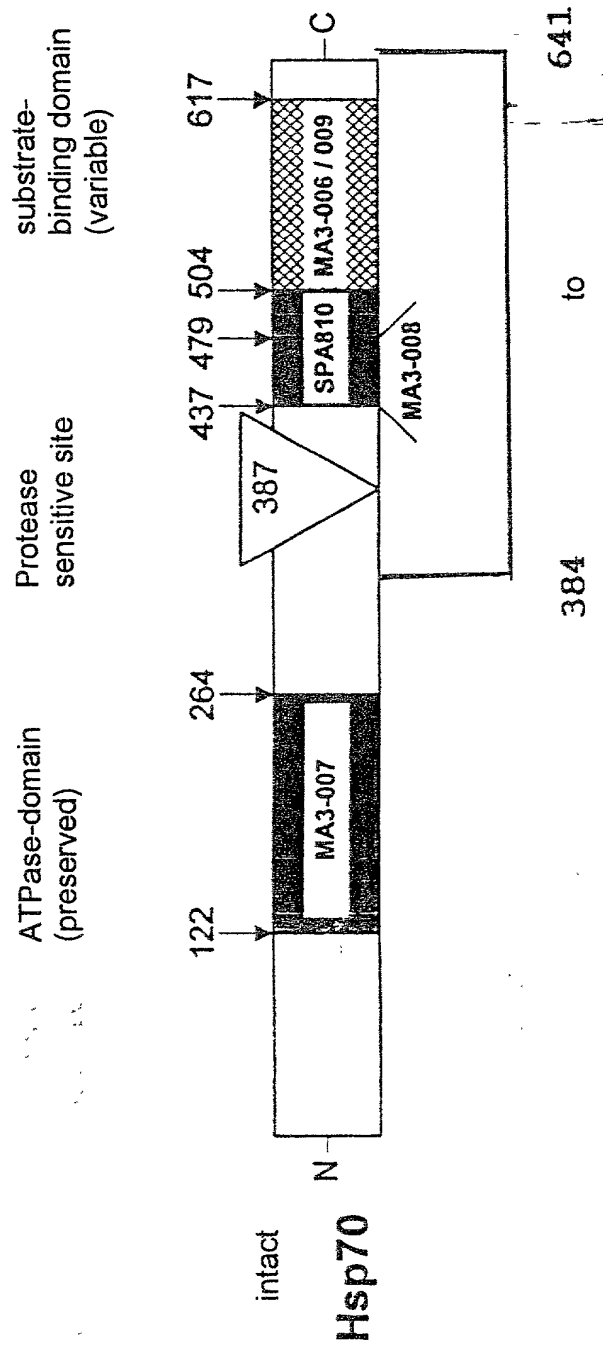


FIG. 5



Intact Hsp70-protein with display of the C-terminal region necessary for the NK-stimulation.



FIG. 6

### 1. Multiple stimulation of NK-cells

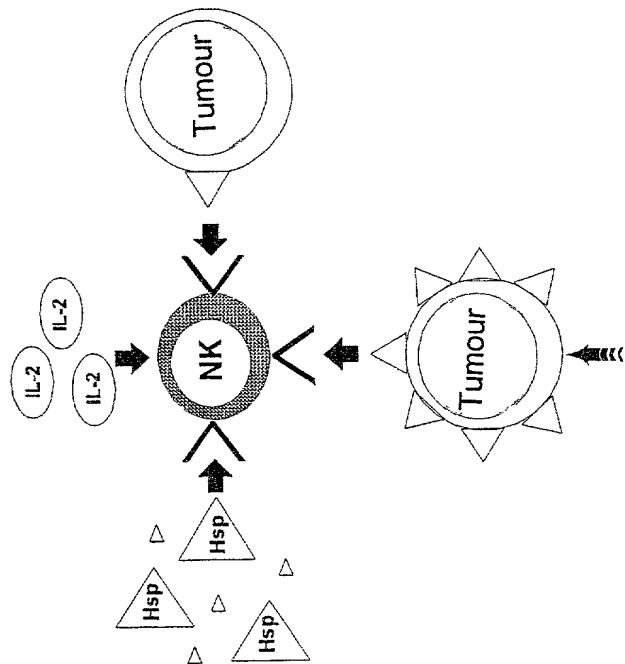
Inactive effector cells (NK- but also T-cells) are multiply stimulated by:

- *ENKASTIM* or derived HSP subunits (here shown as triangles)
- Interleukin-2 (IL-2)
- HSP-expressing tumour cells

As a result, the growth and the killer activity of tumour specific NK- and T-cells is stimulated.

### 2. Increase in the HSP expression on tumour cells by:

- ether lipid treatment
- newly developed substances



# DECLARATION FOR PATENT APPLICATION SOLE INVENTOR

As the named inventor, I hereby declare:

My residence, post office address and citizenship are as stated next to my name below:

I believe that I am the original, first and sole inventor of the subject matter which is claimed and for which a patent is sought on the invention entitled New Use of Hsp70 Protein, described in the specification filed in the U.S. Patent and Trademark Office on 22 September 2000 and assigned serial no. 09/646,835.

I hereby state that I have reviewed and understand the contents of the above-identified specification, including the claims, as amended by any amendment referred to above.

I acknowledge the duty to disclose information which is material to the examination of this application in accordance with Title 37, Code of Federal Regulation, §1.56.

## Prior Foreign Applications

I hereby claim priority benefits under Title 35, United States Code, ' 119 of any foreign application(s) for patent or inventor's certificate listed below. I have also identified below any foreign application(s) for patent or inventor's certificate having a filing date before that of the application on which priority is based.

Country	Application Number	Date of Filing (day, month, year)	Date of Issue (day, month, year)	Priority Claimed Under 35 U.S.C. ' 119
WIPO	PCT/EP99/02165 ✓	29 March 1999 ✓		Yes X No
Germany	DE 198 13 760.5 ✓	27 March 1998 ✓		Yes X No
WIPO	PCT/EP99/02056 ✓	26 March 1999 ✓		Yes X No ✓

## Power of Attorney

And I hereby appoint, both jointly and severally, as my attorneys with full power of substitution and revocation, to prosecute the instant application and to transact all business in the U.S. Patent and Trademark Office connected herewith the following attorneys, their registration numbers being listed after their names:

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Rev. Date 5/99

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I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information knowledge and belief are believed to be true; and further that the statements were made with the knowledge that false statements made willfully are punishable by fine, imprisonment, or both a fine and imprisonment under Section 1001 of Title 18 of the United States; and further that false statements made willfully may jeopardize the validity of any patent issuing on an application in which the false statements were made.

Signature X Dr. Heerl R. Gabriele Date X 28.11.00

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